Appl. No.

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polynucleotide, OC-2 having the amino acid sequence of SEQ ID NO:2, and a cell line transformed with said vector.

- 4. **(AMENDED TWICE)** A pharmaceutical composition comprising an element selected from the group consisting of: a polynucleotide encoding OC-3, a vector comprising said polynucleotide, OC-3 having the amino acid sequence of SEQ ID NO: 3, and a cell line transformed with said vector.
- 7. (AMENDED TWICE) A method for the prevention and or for the treatment of type 1 or type 2 diabetes or of disorders linked to diabetes, for the prevention and or for the treatment of cancer and for the prevention and for the treatment of Waardenburg syndrome, comprising;

administration of a pharmaceutical composition in an amount effective to prevent or reduce the symptoms of diabetes, cancer, and or Waardenburg syndrome, wherein said pharmaceutical composition comprises an element selected from the group consisting of: a polynucleotide encoding a protein of the ONECUT family, a vector comprising said polynucleotide, the polypeptide encoded by said polynucleotide, and a cell line transformed with said vector.

REMARKS

Claims 1, 4, and 7 have been amended, and Claims 2 and 3 have been cancelled. An abstract of the disclosure has been provided on a separate sheet of paper, as required by the Examiner. Changes made to the claims can be seen on a separate page entitled VERSION WITH MARKINGS TO SHOW CHANGES MADE following the signature page. Deletions are in **[bold and brackets]**, and insertions are <u>underlined</u>.

Claim Rejection Under 35 U.S.C. § 112, First Paragraph

Claims 1-9 were rejected under 35 U.S.C. § 112, first paragraph, as containing subject matter which was not described in the specification in such a way as to enable one skilled in the art to which it pertains to make and or use the invention. The Examiner asserts that the specification is enabling for a polynucleotide encoding HNF-6 or OC-2 or OC-3, but that no pharmaceutical property is shown and or enabled. The Examiner further asserts that Claims 7-9 drawn to a method of using the polynucleotide of Claim 1 is not disclosed in the specification.